

K831857 SURGITEK PERCUTANEOUS ARNEGRADE SILITEKAug 11, 1983
63 days to decisionK831857 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k831857/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jun 9, 1983
Decision date	Aug 11, 1983
Days to decision	63 days
Third-party review	No

APPLICANT

Company	Medical Engineering Corp.
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993

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Device record: <https://www.510kdatabase.net/k831857/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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